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Bombshell: Pfizer's "Secret" 2021 Declassified Report. CALL FOR IMMEDIATE WITHDRAWAL OF COVID "VACCINE"

Strategic Analysis and Comprehensive Evaluation of Post-Authorization Adverse Event Data

January 2026

1. Executive Summary: The Institutional Crisis and Data Revelation

The release and subsequent declassification of Pfizer's internal safety data have precipitated a significant reassessment of the global mRNA vaccination campaign. This report, primarily based on the analysis of Professor Michel Chossudovsky, examines the "Cumulative Analysis of Post-Authorization Adverse Event Reports of PF-07302048 (BNT162B2) Received Through 28-Feb-2021". This document, originally classified, was obtained via a Freedom of Information (FOI) procedure in October 2021 and contains data that directly contradicts the early public safety narratives promoted by health organizations.

The core findings of this analysis reveal that within the first two and a half months of the vaccine's rollout—from mid-December 2020 to the end of February 2021—Pfizer recorded 1,223 fatal outcomes and tens of thousands of adverse events. Despite these internal findings being available to the Food and Drug Administration (FDA), the Centers for Disease Control (CDC), and the National Institute of Allergy and Infectious Diseases (NIAID), the vaccine project continued to be endorsed and mandated globally.

Furthermore, the data suggests a widespread and systematic failure of regulatory oversight, culminating in a 2025 FDA order for Pfizer and Moderna to finally issue warnings regarding high risks of heart damage, a risk that was documented in internal reports as early as March 2021. This report calls for the immediate withdrawal of the mRNA vaccine, characterizing the continued marketing of the product as a criminal undertaking and a violation of public trust.

2. Clinical and Regulatory Context: The 2020–2021 Launch Phase

The Pfizer-BioNTech mRNA vaccine (BNT162b2) received its Emergency Use Authorization (EUA) in the United States on December 11, 2020, with the official launch commencing on December 14, 2020. This authorization was granted during a period of intense global pressure from both political entities and pharmaceutical interests, collectively referred to in the sources as "Big Pharma".

The sources assert that as early as March 2021, the U.S. Department of Health and Human Services (HHS) and its subsidiary agencies, including the CDC and the NIAID under Dr. Anthony Fauci, were aware of the underlying dangers associated with the mRNA platform. Despite this awareness, these organizations, in coordination with the World Health Organization (WHO), proceeded to greenlight the worldwide marketing of the vaccine.

Pharmacovigilance Obligations and the BioNTech Partnership Pfizer was responsible for managing post-authorization safety data on behalf of the Marketing Authorization Holder (MAH), BioNTech, in accordance with a predefined Pharmacovigilance Agreement. This agreement necessitated the collection and tabulation of all reported adverse events globally. The resulting integrated analysis (the "Pfizer Report") was designed to provide a comprehensive look at the vaccine's safety profile during its initial phase of distribution.

3. Analysis of Post-Authorization Safety Data (Dec 2020 – Feb 2021)

The declassified Pfizer report provides a granular look at the initial 90 days of the vaccine rollout. This period is critical because it represents the first large-scale real-world application of mRNA technology outside of clinical trials.

3.1 Volume and Geographic Distribution of Reports Between December 1, 2020, and February 28, 2021, a significant but redacted number of doses were shipped globally. During this interval, Pfizer received 42,086 cumulative case reports. These reports were divided into:

- **Medically Confirmed Cases: 25,379 reports.**
- **Non-Medically Confirmed Cases: 16,707 reports.**

These cases contained a total of 158,893 individual adverse events (AEs). The sources highlight that most cases (34,762) were concentrated in a specific group of nations, which may reflect both the speed of the rollout and the robustness of local reporting systems:

- **United States: 13,739 cases.**
- **United Kingdom: 13,404 cases.**
- **Italy: 2,578 cases.**
- **Germany: 1,913 cases.**
- **France: 1,506 cases.**
- **Portugal: 866 cases.**
- **Spain: 756 cases.**

An additional 7,324 cases were distributed across 56 other countries. It is important to note, as the sources emphasize, that these reports were submitted voluntarily, and the magnitude of underreporting is unknown, suggesting the actual numbers could be significantly higher.

3.2 Demographics of Reported Cases The demographic data contained in the report's Table 1 reveals notable disparities in reporting based on gender and age.

- **Gender: There were 29,914 cases reported in females, compared to 9,182 cases in males. Data was not provided for 2,990 cases.**
- **Age Range: The age of affected individuals ranged from 0.01 years to 107 years, with a mean age of 50.9 years.**
- **Pediatric Cases: While the vaccine was initially targeted at adults, 175 cases were reported in individuals aged 17 and younger, including 46 cases in children under 16 and 34 cases in children under 12.**

3.3 Outcomes and Morbidity The outcomes for the 42,086 reported cases as of February 28, 2021, paint a concerning picture of the vaccine's impact on public health.

- **Fatal Outcomes: 1,223 cases resulted in death.**
- **Non-Recovered: 11,361 individuals had not recovered from their adverse event at the time the report was compiled.**
- **Recovered with Sequelae: 520 individuals recovered but were left with permanent or long-term conditions.**
- **Unknown Status: 9,400 cases had unknown outcomes.**
- **Recovered/Recovering: 19,582 cases.**

4. System Organ Class (SOC) Deep Dive: Patterns of Injury

The Pfizer report categorized the 158,893 adverse events by System Organ Class (SOC), identifying the physiological systems most frequently impacted by the injection. The sources highlight several categories that contained more than 2% of the total events in the dataset.

4.1 General Disorders and Administration Site Conditions This was the largest category, accounting for 51,335 total adverse events. Key events within this class included:

- **Pyrexia (Fever): 7,666 instances (18.2% of total cases).**
- **Fatigue: 7,338 instances (17.4% of total cases).**
- **Chills: 5,514 instances (13.1% of total cases).**
- **Vaccination Site Pain: 5,181 instances (12.3% of total cases).**

4.2 Nervous System Disorders There were 25,957 reported events related to the nervous system, making it the second most impacted physiological system after general disorders.

4.3 Musculoskeletal and Connective Tissue Disorders A total of 17,283 events were reported in this category, indicating significant impacts on the structural and functional tissues of the body.

4.4 Gastrointestinal (GI) Disorders GI disorders accounted for 14,096 events. Specific symptoms included:

- **Nausea: 5,182 instances (12.3% of total cases).**
- **Diarrhea: 1,880 instances (4.5% of total cases).**
- **Vomiting: 1,698 instances (4.0% of total cases).**

4.5 Other Significant SOC Categories

- **Respiratory, Thoracic, and Mediastinal Disorders: 8,848 events.**
- **Skin and Subcutaneous Tissue Disorders: 8,476 events.**
- **Injury, Poisoning, and Procedural Complications: 5,590 events.**
- **Infections and Infestations: 4,610 events.**
- **Investigations: 3,693 events.**
- **Blood and Lymphatic System Disorders: This included 1,972 cases of lymphadenopathy (4.7% of cases).**

5. Adverse Events of Special Interest (AESIs): Cardiovascular and Reproductive Health

The Pfizer report included a specific evaluation of Adverse Events of Special Interest (AESIs), which are conditions identified as needing rigorous monitoring due to their severity.

5.1 Cardiovascular AESIs The sources highlight Table 7 of the report, which documents 1,403 cardiovascular cases, representing 3.3% of the total post-marketing dataset. Of these, 241 were medically confirmed. The specific conditions reported included:

- **Tachycardia: 1,098 cases (2.6% of cases).**
- **Arrhythmia: 102 cases.**
- **Myocardial Infarction: 89 cases.**
- **Cardiac Failure: 80 cases.**
- **Acute Myocardial Infarction: 41 cases.**
- **Acute Cardiac Failure: 11 cases.**
- **Cardiogenic Shock and Postural Orthostatic Tachycardia Syndrome: 7 cases each.**
- **Coronary Artery Disease: 6 cases.**

Crucially, the onset latency for these cardiovascular events was extremely short. The range was from less than 24 hours to 21 days, with a median onset of less than 24 hours after the vaccine was administered. This suggests a direct and immediate physiological reaction to the mRNA injection.

5.2 Reproductive Health and Pregnancy One of the most concerning findings highlighted by the sources involves the vaccine's impact on pregnancy. Out of 270 reported pregnancies within the initial cohort, the report documented 23 cases of spontaneous abortions. This data was reportedly available to health authorities while they were still recommending the vaccine to pregnant women.

6. Institutional Failures and the Regulatory Timeline

The analysis provided in the sources suggests a severe disconnect between the safety data known to health agencies and the public health policies enacted between 2021 and 2025.

6.1 Knowledge vs. Action The sources argue that the FDA, CDC, and NIAID were "fully aware" in early March 2021 that the vaccine would lead to an "upward wave of mortality and morbidity". Despite this, the vaccine was "fully endorsed and imposed, nationally and internationally". The FDA reportedly fought to keep this specific document hidden from the public for 55 years, a move described by the sources as a "bombshell" revelation of the agency's complicity.

6.2 The 2025 FDA Warning In August 2025, the FDA finally ordered Pfizer and Moderna to issue formal warnings regarding a "high risk of heart damage," specifically myocarditis, among young men. The sources characterize this as a "bitter irony," noting that this "deadly risk" was documented in the company's own records over four years prior to the official warning.

6.3 Media Disinformation and Conflict of Interest The sources critique the role of mainstream media and "Fact Checkers" in suppressing the findings of the declassified report. A specific conflict of interest is alleged regarding James C. Smith, the former CEO of Reuters and Chairman of the Reuters "Fact Checker" board, who is identified as a top investor and board member of Pfizer. The sources suggest that such overlaps in leadership contributed to the report being "casually ignored" by the media.

7. Corporate Integrity and Historical Precedent

To contextualize the current situation, the sources point to Pfizer's historical record with the U.S. justice system.

7.1 The 2009 Medical Fraud Settlement In 2009, Pfizer reached a $2.3 billion settlement with the U.S. Department of Justice (DOJ) on charges of "fraudulent marketing". This was, at the time, one of the largest healthcare fraud settlements in history.

7.2 The Corporate Integrity Agreement As part of the 2009 settlement, Pfizer entered into an "expansive corporate integrity agreement" with the Office of Inspector General of the HHS. This agreement was designed to put procedures in place to detect and avoid criminal activities similar to those that led to the fraud charges. However, the sources argue that the DOJ's decision to put Pfizer on "probation" rather than pursuing more aggressive criminal charges against executives was a mistake, suggesting that the company's current conduct regarding the mRNA vaccine is a continuation of its past "criminal activities".

8. Conclusion: Ethical Implications and Recommendations

The declassified 2021 Pfizer report provides what the sources describe as a "de facto Mea Culpa". Because the data originates from Pfizer's own internal records, it is considered irrefutable evidence of the product's

risks. The analysis concludes that the mRNA vaccine project has shifted from a public health initiative into a "multibillion dollar operation" that prioritizes business interests over human lives.

Strategic Recommendations from the Sources:

1. **Immediate Withdrawal: The report calls for the immediate and worldwide withdrawal of all Covid mRNA vaccines.**
2. **Cancellation of Mandates: All policies imposing the vaccine nationally or internationally should be canceled.**
3. **Legal Proceedings: The evidence in the confidential report should be used to formulate legal procedures against "Big Pharma," governments, the WHO, and the media.**
4. **Public Disclosure: Full transparency is required regarding all remaining classified safety data held by the FDA and other health agencies.**

The analysis by Professor Chossudovsky frames this as a "Global Coup d'État Against Humanity" and a "criminal undertaking," urging the public and legal bodies to act on the data provided in the company's own 38-page report and its extensive appendices.

Analogy for Understanding: Think of the 2021 Pfizer Report as a "black box" flight recorder found after a series of aviation incidents. If the airline and the aviation regulators had access to the data in that box—showing engine failures and structural flaws—within the first few months of a new plane's operation, but chose to continue flying those planes and forcing passengers to board them for another five years, the eventual release of that "black box" data would represent the ultimate evidence of a systemic failure in safety and ethics. This report is presented as the "black box" of the mRNA vaccine rollout.